Supplementary Online Content

Taichman DB, McGoon MD, Harhay MO, Archer-Chicko C, Sager JS, Murugappan M, Chakinali MM, Palevsky HI, Gallop R. Wide variation in clinicians' assessment of New York Heart Association/World Health Organization functional class in patients with pulmonary arterial hypertension. *Mayo Clin Proc.* 2009;84(7):586-592.

eAPPENDIX. Survey Presented to Physicians and Nurses Regarding the Assessment of Patients' New York Heart Association/World Health Organization Functional Classes

This supplementary material has been peer reviewed, edited, and approved by the authors.

eAPPENDIX. Survey Presented to Physicians and Nurses Regarding the Assessment of Patients'
New York Heart Association/World Health Organization Functional Classes

Dear Colleague,

Please read each of the following brief patient descriptions. Then, please assign a New York Heart Association / World Health Association (NYHA/WHO) class <u>as you would in your own practice</u>.

There are no intended "correct" answers. The goal is simply to see if caregivers come to the same conclusions.

Therefore, <u>please do not discuss</u> these questions with others. Please rank patients as you would in your own practice.

As an incentive and thanks, you will be entered into a raffle to win free registration for the next Pulmonary Hypertension Association International meeting.

Please enter your name and email address on this sheet, DETACH IT from the questionnaire, and turn in this name sheet and your completed questionnaire in the separate boxes near the entry to the conference room. THIS WILL KEEP YOUR ANSWERS ANONYMOUS.

This research has not received industry funding.

Thank you.		
Darren Taichman, MD, PhD		
YOUR NAME	 	
EMAIL ADDRESS		

PATIENTS

1.	38-year-old w	38-year-old woman with idiopathic pulmonary arterial hypertension diagnosed 10 months ago.				
	She demonstrated acute vasoreactivity to inhaled nitric oxide, normalization of her mean					
	pulmonary ar	tery pressure and a 30%	reduction in her puln	nonary vascular resistance. S	She has	
	been treated	with oral calcium channe	el blockers. She work	s as a copy editor.		
NYHA/WHO:	Class I	Class II	Class III	Class IV		
2.	years ago. S warfarin, and	She is being treated with uses supplemental oxyg	a single oral agent for gen. She reports dys	y arterial hypertension diagno pulmonary arterial hypertens onea carrying groceries into h ma. Her 6-minute walk distar	sion, ner	
NYHA/WHO:	Class I	Class II	Class III	Class IV		
3.	dyspnea on e light-headedr pressure is 1	exertion. She becomes s	short of breath after cledema. At right heart ary artery pressure 49	rtension and 8 months progre imbing two flights of stairs. S catheterization, her right atria mm Hg, cardiac index 1.6 La	She denies	
NYHA/WHO:	Class I	Class II	Class III	Class IV		
4.	hypertension	. She also uses supplen	nental oxygen at 6 lite	HIV associated pulmonary a ers per minute. At rest her O_2 and talking on the phone.		
NYHA/WHO:	Class I	Class II	Class III	Class IV		
5.	mmHg was n transplantation mm Hg, and and oral silde	noted at a catheterization on. The right atrial press cardiac index 2.9 L/min/r	performed one year a cure was 8 mm Hg, pu m ² . Since then, she h comes light-headed a	pulmonary artery pressure of ago during evaluation for pote Imonary artery occlusion pres as been treated with inhaled fter climbing a flight of stairs,	ential liver ssure 10 iloprost	
NVHA/WHO:	Class I	Class II	Class III	Class IV		

	following evalu	ation for a syncopal e	episode one week ago	. She is relatively sedentary at basel	ine
	and denies any	recent change in he	r activities due to sho	rtness of breath. She performs activit	ies
	of daily living w	rithout noted significa	nt shortness of breath	. On echo, her right heart is dilated.	At
	cardiac cathete	erization the right atria	al pressure 10 mm Hg	, mean pulmonary artery pressure 45	
	mm Hg, cardia	c index 2.1L/min/m² a	and pulmonary artery	occlusions pressure is 7 mm Hg.	
NYHA/WHO:	Class I	_ Class II	Class III	Class IV	
7.	A 49-year-old v	voman with sclerode	rma associated pulmo	nary arterial hypertension, diagnosed	
	three years ago	o. She has been trea	ted with single agent	for pulmonary arterial hypertension,	
	warfarin and ox	cygen. She reports s	hortness of breath car	rying groceries. She denies	
	lightheadness	or leg swelling. Her s	six-minute walk distan	ce is 390 meters.	
NYHA/WHO:	Class I	Class II	Class III	Class IV	
8.	51-year-old wo	man with anorectic a	gent associated pulmo	onary arterial hypertension who has	
	been treated with subcutaneous treprostinil for the last 18 months. She has site pain that				
	requires narcotics for relief on a daily basis. She is a kindergarten teacher and continues to work				
	full time. She r	notes dyspnea when	playing games outdoo	ors with her students.	
NYHA/WHO:	Class I	Class II	Class III	Class IV	
9.	56-year-old ma	ın with idiopathic pulr	monary arterial hyperte	ension, treated with intravenous	
	epoprostenol for the last year. He does clerical work. He denies shortness of breath. He does				
	feel tired at the end of a long workday. He notes mild bilateral lower extremity edema. A recent				
	cardiac catheterization revealed a right atrial pressure of 11 mm Hg, mean pulmonary artery				
	pressure of 40 mm Hg, cardiac index of 2 L/min/m ² and a pulmonary artery occlusion pressure of				
	9 mm Hg.				
NYHA/WHO:	Class I	Class II	Class III	Class IV	
10.	37-year-old wo	man with scleroderm	a associated pulmona	ary arterial hypertension. She was no	t
	responsive to inhaled nitric oxide at cardiac catheterization one year ago and has since been				
	treated with ora	al bosentan. She cur	rently denies limitation	ns due to dyspnea or fatigue. She wo	rks

35-year-old woman with scleroderma associated pulmonary arterial hypertension, diagnosed

6.

as a receptionist.

NYHA/WHO:	Class I	Class I	l	Class III	Class IV
<u>PLEA</u>	SE COMPLE	TE THE FOLLOW	ING INFO	<u>RMATION</u>	
Your p	·	k all that apply):			
	•	n 🖵 Nurse	∟ Nur	se Practitioner	☐ Research Coordinator
Your i	nvolvement w	vith patients with pu	ılmonary a	rterial hypertensi	on (check all that apply):
		are 🖵 coord			
		which you practice			
Your p	orimary practi	ce specialty:			
	upulmona 🖵	ry 🖵 cardiolo	gy [rheumatology	
	internal i	medicine without s	ubspecialty	v □ critical	care
	☐ other				
Your p	oractice settin	g:			
	🖵 private c	ffice-based practic	е 🖵	academic medic	al center
	☐ private h	ospital-based prac	tice	☐ other	
Numb	er of years yo	ou have been treati	ng patients	s with pulmonary	hypertension:
	🗖 0 to less	than 1 year	☐ 1 to less	than 3 years	$lue{}$ 3 to less than 5 years
	☐ 5 to less	than 10 years	☐ 10 or	more years	
Please year:	e estimate the	e number of new pa	atients with	ı pulmonary arter	ial hypertension you see each
	□ <10	□ 10-20	1 21-50	□>50	
-	u currently us ng therapy?	se the WHO or NYF	HA function	al classification a	as part of your assessment when
	☐ Yes	☐ No			

Have you enrolled patients into clinical trials or studies of pulmonary arterial hypertension?

	☐ Yes	☐ No			
Please	check all pulr	monary arterial hyper	tension therapies th	nat you have prescribe	ed or managed:
	acalcium cl	hannel blockers	☐ bosentan	lacksquare ambrisentan	☐ sitaxsentan
	🖵 sildenafil	☐ tedalafil	☐ inhaled iloprost	intravenous	treprostinil
	☐ subcutaneous treprostinil ☐ intravenous epoprostenol				
	(I am not	directly involved in pa	atients' clinical care)	
Do you think the patients described in this questionnaire resemble patients you might see in your own practice?					
	☐ Yes	☐ No			
	check all iten ny as apply):	ns that you consider v	when determining a	patient's NYHA/WHC) class (check
	☐ symptoms	s 🖵 medications	s used 🖵 hem	nodynamic values	
☐ physical examination ☐ 6 minute walk distance					
	echocardi	iographic findings	patients' lifest	yle 🖵 patient's o	occupation
☐ Requirements to get insurance approval for the therapy I think best for the patient					
	☐ other				

THANK YOU!